U.S. Patent and Trademark Office,
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless i

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
( Not for submission under 37 CFR 1.99)

Application Number		10723370	
Filing Date		2003-11-26	
First Named Inventor Gros		5	
Art Unit		2626	
Examiner Name	Lamo	ont Spooner	
Attorney Docket Number		JNG 98001C	

					U.S.I	PATENTS			Remove		
Examiner Initial*	Cite No	Patent Number	Kind Code <sup>1</sup>	Issue D	late	Name of Patentee or Applicant of cited Document			Pages,Columns,Lines w Relevant Passages or R Figures Appear		
	1	5280573		1994-01	1-18	Kuga et al.					
	2	5576755		1996-11	1996-11-19 Davis						
	3	5757417		1998-05	5-26	Aras et al.					
	4	6075550		2000-06	5-13	Lapierre					
If you wis	h to a	dd additional U.S. Pater	nt citatio	n inform	ation pl	ease click the	Add button.		Add		
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove		
Examiner Initial*	Cite No	Publication Number	Kind Code <sup>1</sup>	Publica Date	tion	Name of Patentee or Applicant		Releva		Lines where ges or Relev	
	1										
If you wis	h to a	dd additional U.S. Publi	shed Ap	plication	citation	n information p	please click the Ad	d button	Add		_
				FOREIG	SN PAT	ENT DOCUM	IENTS		Remove		_
Examiner Initial*	Cite No	Foreign Document Number <sup>3</sup>	Country Code <sup>2</sup>		Kind Code4	Publication Date	Name of Patente Applicant of cited Document	e or V	vhere Rel	or Relevant	TS

	Application Number		10723370	
NEODWATION DIGGI COURT	Filing Date		2003-11-26	
NFORMATION DISCLOSURE STATEMENT BY APPLICANT	First Named Inventor Gros		SS	
Not for submission under 37 CFR 1.99)	Art Unit		2626	
Not for Submission and or or it 1.557	Examiner Name Lamo		amont Spooner	
	Attorney Docket Number		JNG 98001C	

	1											
If you wis	h to a	dd addit	ional Foreign	Patent Doo	cument cita	ition in	nformation pl	ease click	the Add butto	n Add	1	
				NON	-PATENT	LITER	RATURE DO	CUMENT	S	Remove		
Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item Initials* No look, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.						Тв						
	1											
If you wis	h to a	dd addit	ional non-pai	tent literatur	e documer	nt citat	tion informat	ion please	click the Add	button A	dd	
					EXAM	IINER	SIGNATUR	E				
Examiner	Signa	ture						Date	Considered			
*EXAMIN	ER: Ir	itial if re	ference con:	sidered, who	ether or no	t citati	ion is in conf	ormance v	with MPEP 609	. Draw line	through a	

See Kind Codes of USPTO Plantin Documents at twee\_USPTO\_EDUC on MPP 901.04. \* Enter office and assess the document, by the to-electr code (WIPO Standard ST3.). \*\*Enter office the Separanee petral comments, the reduction to the parent for sering role the Engerer must proceed the serial number of the patent counter.

\*\*Kind of Cocument by the appropriate symbols as indicated on the document under WIPO Standard ST.16 if possible. \*\*Applicant is to place a check mark here it from the patent counter.

\*\*Comment of the Parent Comment of the Parent Co

citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

# INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10723370		
Filing Date		2003-11-26		
First Named Inventor	Gros	Gross		
Art Unit		2626		
Examiner Name	Lamo	nt Spooner		
Attorney Docket Number		JNG 98001C		

#### CERTIFICATION STATEMENT

Please see 37 CFF	1.97 and 1.98	to make the appropr	riate selection(s):
-------------------	---------------	---------------------	---------------------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the information disclosure statement. See 3.7 CFR 1.97(eVI.)

### OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no tem of information contained in the information disclosure statement was known to [2] any individual designated in 37 CFR 1.58(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.59(c) the statement of the filing of the information disclosure statement. See 37 CFR 1.59(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 1.59(c) and the statement of the

See attached certification statement.

Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

## None

### SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/John N. Gross/	Date (YYYY-MM-DD)	2006-08-04
Name/Print	John N. Gross	Registration Number	34175

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life and by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C.12 Card 37 CFR.

1.14. This collection is estimated to take I hour to complete, including gathering, preparing and submitting the completed application from to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenar's Office, and Superiment of Commence, P. 0. Bot 1450, Alexandria, V.32.211-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P. 0. Box 1450, Alexandria, V.32.231-1450.

### Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that (1) the general authority for the collection of this information is \$3 U.S.C. 2(b)(2); (2) famishing of the information solicided is voluntary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan Kollie is to process and/or examine your submission related to a patient application or patient. If you do not furnish the requested process and/or examine your submission related to a patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested related to the patient application or patient. If you do not furnish the requested to the patient process and/or examine your submission, which may related that the patient process and/or examine your submission, which may

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
  - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement neodiations.
  - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record partains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
  - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974. as amended, pursuant to 5 U.S.C. 552a(m.).
  - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
    may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
    to the Patent Cooperation Treaty.
  - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
  - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2504 and 2506. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make relevant for the state of the s
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S. C. 12(b) or issuance of a patient pursuant to 35 U.S. C. 15.1 Further, a record may be disclosed, subject to the limitations of 37 CFR.114, as a routine use, to the public if the record via flori of mapplication which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inseptions or an issued patient.
  - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.